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XX WPI; 2000-431566/37.  
DR N-PSDB; AAA78402.  
XX  
PT Forty seven human nucleic acids encoding secreted proteins, useful in  
PT the treatment, prevention and diagnosis of cancers, disorders of the  
PT immune system, angiogenesis disorders, neurological diseases and  
PT hyperproliferative disorders -  
XX  
PS Claim 11: Page 496; 562pp; English.  
XX  
CC The polynucleotide sequence given in AAA78381 to AAA78432 encode the  
CC human secreted proteins given in AAB24437 to AAB24604. Human secreted  
CC proteins have activities based on the tissues and cells the genes are  
CC expressed in. Examples of activities include: cytostatic; antineoplastic;  
CC antidiabetic; antiinflammatory; ophthalmological; antirheumatic;  
CC antithrombotic; antiproliferative; angiogenic; cardiac; anti-HIV;  
CC neurotropic; neuroprotective; antimicrobial and antiparkinsonian.  
CC Human secreted protein polynucleotides, polypeptides, antagonists and/or  
CC agonists may be useful in treating, preventing, and/or diagnosing other  
CC diseases, disorders, and/or conditions such as: (a) cancers; (b)  
CC disorders of the immune system; (c) angiogenesis disorders; (d)  
CC hyperproliferative disorders; (e) cardiovascular disorders; (f) diseases  
CC associated with increase apoptosis; (g) neurological diseases; and  
CC (h) infectious diseases. They are also used to promote wound healing.  
CC AAA78372 to AAA78380 and AAB24436 represent sequences used in the  
CC exemplification of the present invention.  
XX  
SQ Sequence 191 AA:  
  
Query Match 100.0%; Score 1002; DB 21; Length 191;  
Best Local Similarity 100.0%; Pred. No. 2,2e-111;  
Matches 190; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
  
QY 1 MMNFOPPSKARASOMMTFFILFFPSFGVCTLAITIRLKPSADCGPFGLPFIH 60  
DB 1 MMNFOPPSKARASOMMTFFILFFPSFGVCTLAITIRLKPSADCGPFGLPFIH 60  
QY 61 SIYSWIDTLSTRPGYLMVWVIYRNIGSVHFFILTLIVLITLYLWQITEGKIMIRLL 120  
DB 61 SIYSWIDTLSTRPGYLMVWVIYRNIGSVHFFILTLIVLITLYLWQITEGKIMIRLL 120  
QY 121 HEQIINEGKDKMFLIEKLIRLOMEKKANPSSVLERREVEOGFHLGHDGSLDLRSR 180  
DB 121 HEQIINEGKDKMFLIEKLIRLOMEKKANPSSVLERREVEOGFHLGHDGSLDLRSR 180  
QY 181 RSYOEGNPRA 190  
DB 181 RSYOEGNPRA 190

RESULT 2  
AAB83082  
ID AAB83082 standard; Protein; 191 AA.  
XX  
AC AAB83082;  
XX  
DT 29-JUN-2001 (first entry)  
XX  
DE Human CASB6411-related partial polypeptide #2.  
XX  
KW Human; CASB6411; vaccine; gene therapy; immunoprophylaxis;  
KW ovarian cancer; colon cancer; autoimmune disease.  
XX  
OS Homo sapiens.  
XX  
PN WO200123417-A2.  
XX  
PD 05-APR-2001.  
XX  
PF 27-SEP-2000; 2000MO-EP09500.  
XX  
PR 30-SEP-1999; 99GB-0023154.

PR 07-JUL-2000; 2000GB-0016839.  
XX  
PA (SMIR ) SMITHKLINE BEECHAM BIOLOGICALS.  
XX  
PI Vinals De Bassols YC;  
XX  
DR WPI; 2001-316133/33.  
DR N-PSDB; AAF82463.  
XX  
PT Novel CASB6411 polypeptides useful in diagnostics, and as vaccines for  
PT prophylactic and therapeutic treatment of cancers, particularly ovarian  
PT and colon cancers, autoimmune diseases and related conditions -  
XX  
PS Disclosure; Page 67; 95pp; English.  
XX  
CC The present sequence is provided in a specification relating  
CC to CASB6411 polypeptides comprising a sequence having at least 70%  
CC identity to a sequence of 460 or 154 amino acids fully defined in  
CC the specification. CASB6411 polypeptides and polynucleotides are  
CC useful for treating a subject by immunoprophylaxis or therapy.  
CC The CASB6411 polypeptides are useful in diagnostics, and as  
CC vaccines for prophylactic and therapeutic treatment of cancers,  
CC particularly ovarian and colon cancers, autoimmune diseases and related  
CC conditions. CASB6411 polypeptides are also useful for the  
CC structure-based design of agonists, antagonists or inhibitors of the  
CC polypeptide.  
XX  
SQ Sequence 191 AA:  
  
Query Match 100.0%; Score 1002; DB 22; Length 191;  
Best Local Similarity 100.0%; Pred. No. 2,2e-111;  
Matches 190; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
  
QY 1 MMNFOPPSKARASOMMTFFILFFPSFGVCTLAITIRLKPSADCGPFGLPFIH 60  
DB 2 MMNFOPPSKARASOMMTFFILFFPSFGVCTLAITIRLKPSADCGPFGLPFIH 61  
QY 61 SIYSWIDTLSTRPGYLMVWVIYRNIGSVHFFILTLIVLITLYLWQITEGKIMIRLL 120  
DB 62 SIYSWIDTLSTRPGYLMVWVIYRNIGSVHFFILTLIVLITLYLWQITEGKIMIRLL 121  
QY 121 HEQIINEGKDKMFLIEKLIRLOMEKKANPSSVLERREVEOGFHLGHDGSLDLRSR 180  
DB 122 HEQIINEGKDKMFLIEKLIRLOMEKKANPSSVLERREVEOGFHLGHDGSLDLRSR 181  
QY 181 RSYOEGNPRA 190  
DB 182 RSYOEGNPRA 191

RESULT 3  
AAM79104  
ID AAM79104 standard; Protein; 268 AA.  
XX  
AC AAM79104;  
XX  
DT 06-NOV-2001 (first entry)  
XX  
DE Human protein SEQ ID NO 1766.  
XX  
KW Human; cytokine; cell proliferation; cell differentiation; gene therapy;  
KW vaccine; peptide therapy; stem cell growth factor; haematopoiesis;  
KW tissue growth factor; immunomodulatory; cancer; leukaemia;  
KW nervous system disorder; arthritis; inflammation.  
XX  
OS Homo sapiens.  
XX  
PN WO200157190-A2.  
XX  
PD 09-AUG-2001.  
XX  
PF 05-FEB-2001; 2001WO-US04098.  
XX

PR 03-FEB-2000; 2000US-0496914.  
PR 27-APR-2000; 2000US-0560875.  
PR 20-JUN-2000; 2000US-0598075.  
PR 19-JUL-2000; 2000US-0620325.  
PR 01-SEP-2000; 2000US-0624936.  
PR 15-SEP-2000; 2000US-0663561.  
PR 20-OCT-2000; 2000US-0693325.  
PR 30-NOV-2000; 2000US-0728422.  
XX  
XX (HYSE-) HYSEQ INC.  
PI Tang YT, Liu C, Drmanac RT, Asundi V, Zhou P, Xu C, Cao Y, Ma Y;  
PI Zhao OA, Wang D, Zhang J, Ren F, Chen R, Wang ZW;  
PI Xue AJ, Yang Y, Wejhrman T, Goodrich R;  
XX WPI: 2001-476283/51.  
XX N-PSDB: AAK52237.  
PT Nucleic acids encoding polypeptides with cytokine-like activities,  
PT useful in diagnosis and gene therapy -  
XX  
XX Claim 20; Page 4113-4114; 6221pp; English.  
XX  
XX The invention relates to polynucleotides (AAK51456-AAK53435) and the  
CC encoded polypeptides (AAM78323-AAK80302) that exhibit activity elating to  
CC cytokine, cell proliferation or cell differentiation or which may induce  
CC production of other cytokines in other cell populations. The  
CC polynucleotides and polypeptides are useful in gene therapy, vaccines or  
CC peptide therapy. The polypeptides have various cytokine-like activities,  
CC e.g. stem cell growth factor activity, haematopoiesis regulating  
CC activity, tissue growth factor activity, immunomodulatory activity and  
CC activin/inhibin activity and may be useful in the diagnosis and/or  
CC treatment of cancer, leukemia, nervous system disorders, arthritis and  
CC inflammation.  
CC Note: Records for SEQ ID NO 2110 (AAK52581), 2111 (AAK52582) and 3666  
CC (AAM80020) are omitted as the relevant pages from the sequence listing  
CC were missing at the time of publication.  
XX  
XX Sequence 268 AA;  
SQ  
Query Match 100.0%; Score 1002; DB 22; Length 268;  
Best Local Similarity 100.0%; Pred. No. 3.4e-111;  
Matches 190; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 MANNPPSKARRASQMMFFFLFFPSTGYLCTLATITWRKASACCGPRGRLPRT 60  
DB 79 MANNPPSKARRASQMMFFFLFFPSTGYLCTLATITWRKASACCGPRGRLPRT 138  
QY 61 STYSWIDLTSPGGLVWVWYIRNLIGSVHFFITLVLITTYLYWOITBGRKIMIRL 120  
DB 139 STYSWIDLTSPGGLVWVWYIRNLIGSVHFFITLVLITTYLYWOITBGRKIMIRL 198  
QY 121 HEQIINEGKDKMFLIEKLIKIDMEKKANPSSVLVERREVEQGFHLGHEHDSIDLRSR 180  
DB 199 HEQIINEGKDKMFLIEKLIKIDMEKKANPSSVLVERREVEQGFHLGHEHDSIDLRSR 258  
QY 181 RSVQEGNPR 190  
DB 259 RSVQEGNPR 268  
RESULT 4  
ID ABB11361 standard; peptide: 280 AA.  
XX ABB11361;  
XX 11-JAN-2002 (first entry)  
XX Human IAK-4p homologue, SEQ ID NO:1731.  
DE Human: cytokine; cell proliferation; cell differentiation; growth factor;  
KW haematopoiesis regulation; tissue growth; immunomodulator; activin;  
KW

KW inhibin: chemotaxis; chemokinesis; thrombolysis; oncogenesis;  
KW proliferation; metastasis; cancer; tumour; haematopoietic disorder;  
KW myeloid cell disorder; lymphoid cell disorder; asthma; arthritis;  
KW chronic inflammatory condition; proliferative retinopathy;  
KW atherosclerosis; coronary heart disease; arterial ischaemia;  
KW bone disorder; osteoporosis; vascular growth disorder;  
KW tissue regeneration; wound healing; infection; immune disorder;  
KW cell culture; drug screening; gene therapy; antiinflammatory;  
KW antiasthmatic; antiarthritis; haemostatic; antiarteriosclerosis;  
KW cytosolic; osteopathic; vasotropic; cardiant; virucide; antibacterial;  
KW antifungal; vulnery; antiulcer.  
XX  
XX Homo sapiens.  
XX  
XX WO200157188-A2.  
XX  
XX 09-AUG-2001.  
XX  
XX 05-FEB-2001; 2001WO-US03800.  
XX  
XX 03-FEB-2000; 2000US-0496914.  
XX 27-APR-2000; 2000US-0560875.  
XX  
XX (HYSE-) HYSEQ INC.  
PI Tang YT, Liu C, Drmanac RT;  
XX WPI: 2001-457740/49.  
XX N-PSDB: ABA08605.  
PT Human proteins and DNA encoding sequences useful for preventing,  
PT treating or ameliorating a medical condition in a mammalian subject  
PT e.g. arthritis and cancer -  
XX  
XX Claim 20; Page 173; 1963pp; English.  
PS  
PS Sequences ABB10981-ABB12330 represent 1350 novel human polypeptides, and  
CC sequences ABA08225-ABA09574 represent nucleic acids encoding them. The  
CC invention also relates to vectors and recombinant host cells comprising a  
CC nucleotide of the invention, methods of producing the novel polypeptides,  
CC antibodies against the polypeptides, methods of detecting the nucleotides  
CC or polypeptides in a sample, and methods of identifying compounds which  
CC bind to polypeptides of the invention. Although novel, many of the  
CC polypeptides of the invention have homology to known proteins, thereby  
CC giving an insight into their probable biological activities, and hence  
CC potential therapeutic applications. The polypeptides of the invention may  
CC have various activities, including cytokine, cell proliferation or cell  
CC differentiation activities; stem cell growth factor activity;  
CC haematopoiesis regulatory activity; tissue growth activity;  
CC immunomodulatory activity; activin- or inhibin-related activities;  
CC chemotactic or chemokinetic activities; haemostatic, thrombotic or  
CC thrombolytic activities; receptor or ligand activities; or may be  
CC involved in oncogenesis, cancer cell proliferation or metastasis.  
CC Depending on their biological activities, polypeptides and nucleotides of  
CC the invention are useful for preventing, treating or ameliorating medical  
CC conditions, e.g., by protein or gene therapy. Such conditions include  
CC cancers, haematopoietic disorders (e.g., myeloid or lymphoid cell  
CC disorders), chronic inflammatory conditions (e.g., asthma or arthritis),  
CC proliferative retinopathy, atherosclerosis, coronary heart disease,  
CC arterial ischaemia, bone disorders (e.g., osteoporosis), and abnormal  
CC vascular growth. Polypeptides involved with tissue regeneration and  
CC repair (or nucleic acids encoding them) may be used to promote wound  
CC healing (e.g., of burns, incisions and ulcers), while those with  
CC immunomodulatory activities may be used in the treatment of viral,  
CC bacterial and fungal infections in addition to immune disorders.  
CC Polypeptides with growth factor activity may be used in cell cultures to  
CC promote cell growth. For example, such polypeptides may be used to  
CC manipulate stem cells in culture to give rise to neuroepithelial cells  
CC that can be used to augment or replace cells damaged by illness,  
CC autoimmune disease or accidental damage. The polypeptides and nucleotides  
CC may also be used in the diagnosis of the above conditions, and in drug  
CC screening techniques. The present sequence represents a novel human  
CC polypeptide of the invention.

XX	Sequence	280 AA:	Query Match	100.0%;	Score 1002;	DB 22;	Length 280;
SD			Best Local Similarity	100.0%;	Pred. No. 3,6e-111;		
			Matches 190;	Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0;
QY	1	MNNPQPPSKAMVASOMMTFFILFFPPSTGYGLCTLATITIMKLKPSADCGPRGLPLFIH	60				
Db	91	MNNPQPPSKAMVASOMMTFFILFFPPSTGYGLCTLATITIMKLKPSADCGPRGLPLFIH	150				
QY	61	SIYSWIDLTSTRPGILMVVMIYRNILGSHVFFIITLLIYLITTYLWQITTEGRKIMIRLL	120				
Db	151	SIYSWIDLTSTRPGILMVVMIYRNILGSHVFFIITLLIYLITTYLWQITTEGRKIMIRLL	210				
QY	121	HEQIINEGKDKMFLIEKLKIDQMEKANKPSSLVTERREVEQGGFLHIGEHGSDLDLRSR	180				
Db	211	HEQIINEGKDKMFLIEKLKIDQMEKANKPSSLVTERREVEQGGFLHIGEHGSDLDLRSR	270				
QY	181	RSVQEGNRPRA 190					
Db	271	RSVQEGNRPRA 280					

RESULT_5	
AA080088	
ID	AA080088 standard; Protein; 280 AA.
XX	
AC	AA080088;
XX	
DT	06-NOV-2001 (first entry)
XX	
DE	Human protein SEQ ID NO 3734.
XX	
KW	Human; cytokine; cell proliferation; cell differentiation; gene therapy;
KW	vaccine; peptide therapy; stem cell growth factor; haematopoiesis;
KW	tissue growth factor; immunomodulatory; cancer; leukaemia;
KW	nervous system disorder; arthritis; inflammation.
XX	
OS	Homo sapiens.
XX	
PN	WO200157190-A2.
XX	
PD	09-AUG-2001.
XX	
PF	05-FEB-2001; 2001WO-US04098.
XX	
PR	03-FEB-2000; 2000US-0496914.
PR	27-APR-2000; 2000US-0560875.
PR	20-JUN-2000; 2000US-0598075.
PR	19-JUL-2000; 2000US-0620325.
PR	01-SEP-2000; 2000US-0654936.
PR	15-SEP-2000; 2000US-0663561.
PR	20-OCT-2000; 2000US-0693325.
PR	30-NOV-2000; 2000US-0728422.
XX	
PA	(HYSE-) HYSEQ INC.
XX	
PI	Tang YT, Liu C, Drmanac RT, Asundi V, Zhou P, Xu C, Cao Y, Ma Y;
PI	Xue QA, Wang D, Wang J, Zhang J, Ren F, Chen R, Wang ZW;
PI	Zhao AJ, Yang Y, Wehrman T, Goodrich R;
XX	
DR	WPI: 2001-476283/51.
DR	N-PSDB; AAK53221.
XX	
PT	Nucleic acids encoding polypeptides with cytokine-like activities,
PT	useful in diagnosis and gene therapy -
XX	
PS	Claim 20; Page 421; 6221pp; English.
XX	
CC	The invention relates to polynucleotides (AAK51456-AAK53435) and the
CC	encoded polypeptides (AAK78323-AAK80302) that exhibit activity elating to
CC	cytokine, cell proliferation or cell differentiation or which may induce

production of other cytokines in other cell populations. The polynucleotides and polypeptides are useful in gene therapy, vaccines or peptide therapy. The polypeptides have various cytokine-like activities, e.g. stem cell growth factor activity, haematopoiesis regulating activity, tissue growth factor activity, immunomodulatory activity and activin/inhibin activity and may be useful in the diagnosis and/or treatment of cancer, leukaemia, nervous system disorders, arthritis and inflammation.

Note: Records for SEO ID NO 2110 (AAK52581), 2111 (AAK52582) and 3666 (AAM80020) are omitted as the relevant pages from the sequence listing were missing at the time of publication.

	Query Match	100.0%	Score 1002;	DB 22;	Length 280;
	Best Local Similarity	100.0%	Pred. No. 3.6e-11;		
	Matches 190;	Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0.
QY	1	MNFPQPSKAMRASOMTFFIFLFFPSPTGVCLTAITIMRLKPSADCGPRGLPLFIH	60		
Db	91	MNFPQPSKAMRASOMTFFIFLFFPSPTGVCLTAITIMRLKPSADCGPRGLPLFIH	150		
QY	61	STYSWIDILSTRPGILWVWYIIRNLIGSHVHFFILITLYLITLYLWQITBEKRIIMRIIL	120		
Db	151	STYSWIDILSTRPGILWVWYIIRNLIGSHVHFFILITLYLITLYLWQITBEKRIIMRIIL	210		
QY	121	HEQIINEGKDKKFLLEKLIKILQDMKKKAPSSLYLERRREVOQGFHLHLEHGSLDLRSR	180		
Db	211	HEQIINEGKDKKFLLEKLIKILQDMKKKAPSSLYLERRREVOQGFHLHLEHGSLDLRSR	270		
QY	181	RSVOEGCNPRA 190			
Db	271	RSVOEGCNPRA 280			

```

RESULT 6
AAB95481
ID AAB95481 standard; Protein; 330 AA.
XX
XX AAB95481;
AC
XX 26-JUN-2001 (first entry)
DT
XX
XX Human protein sequence SEQ ID NO:18002.
DE
XX
XX Human; primer: detection; diagnosis; antisense therapy; gene therapy.
XX
XX Homo sapiens.
OS
XX EP1074617-A2.
PN
XX
XX 07-FEB-2001.
PD
XX
XX 28-JUL-2000; 2000EP-0116126.
PF
XX
XX 29-JUL-1999; 99JP-0248036.
PR 27-AUG-1999; 99JP-0300253.
PR 11-JAN-2000; 2000JP-0118776.
PR 02-MAY-2000; 2000JP-0183767.
PR 09-JUN-2000; 2000JP-0241899.
XX
XX
XX (HELI-) HELIX RES INST.
PA
XX
XX Ota T, Isogai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;
PI Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;
XX
XX WPI; 2001-318749/34.
DR
XX
XX Primer sets for synthesizing polynucleotides, particularly the 5602
PT full-length cDNAs defined in the specification, and for the detection
PT and/or diagnosis of the abnormality of the proteins encoded by the
PT full-length cDNAs -
XX
XX

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Db      198  MNFOPPSKAMRASQMMTEFFILFFPSPFTGVCTLAITITWRLKPSADCGPFGDLPLFIH 257
QY      61  SIYSWIDTSTRGYLWVWYIYNLIGSVHFFFLITLIVLITITLYWQTEGKRIIMRL 120
Db      258  SIYSWIDTSTRGYLWVWYIYNLIGSVHFFFLITLIVLITITLYWQTEGKRIIMRL 317
QY      121  HEQIINEGKDKMFLIEKLKIQDMEKANPSSVLERREVEQOGFLHGHDSGLDLRSR 180
Db      318  HEQIINEGKDKMFLIEKLKIQDMEKANPSSVLERREVEQOGFLHGHDSGLDLRSR 377
QY      181  RSVQEGNPRA 190
Db      378  RSVQEGNPRA 387

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# RESULT 8

AAB83081 standard; Protein; 438 AA.

AAB83081;

29-JUN-2001 (first entry)

Human CASB6411-related partial polypeptide #1.

Human: CASB6411; vaccine; gene therapy; immunoprophylaxis;  
ovarian cancer; colon cancer; autoimmune disease.

Homo sapiens.

WO200123417-A2.

05-APR-2001.

27-SEP-2000; 2000WO-EP09500.

30-SEP-1999; 99GB-0023154.

07-JUL-2000; 2000GB-0016839.

(SMIK ) SMITHKLINE BEECHAM BIOLOGICALS.

Vinals De Bassols YC;

WPI: 2001-316133/73.

N-PSDB; AAF82462.

Novel CASB6411 polypeptides useful in diagnostics, and as vaccines for  
prophylactic and therapeutic treatment of cancers, particularly ovarian  
and colon cancers, autoimmune diseases and related conditions -  
Disclosure; Page 66; 95pp; English.

The present sequence is provided in a specification relating  
to CASB6411 polypeptides comprising a sequence having at least 70%  
identity to a sequence of 460 or 154 amino acids fully defined in  
the specification. CASB6411 polypeptides and polynucleotides are  
useful for treating a subject by immunoprophylaxis or therapy.  
The CASB6411 polypeptides are useful in diagnostics, and as  
vaccines for prophylactic and therapeutic treatment of cancers,  
particularly ovarian and colon cancers, autoimmune diseases and related  
conditions. CASB6411 polypeptides are also useful for the  
structure-based design of agonists, antagonists or inhibitors of the  
polypeptide.

Sequence 438 AA;

Query Match 100.0%; Score 1002; DB 22; Length 438;  
Best Local Similarity 100.0%; Pred. No. 6-111;  
Matches 190; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1 MNFOPPSKAMRASQMMTEFFILFFPSPFTGVCTLAITITWRLKPSADCGPFGDLPLFIH 60  
|||||

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Db      249  MNFOPPSKAMRASQMMTEFFILFFPSPFTGVCTLAITITWRLKPSADCGPFGDLPLFIH 308
QY      61  SIYSWIDTSTRGYLWVWYIYNLIGSVHFFFLITLIVLITITLYWQTEGKRIIMRL 120
Db      309  SIYSWIDTSTRGYLWVWYIYNLIGSVHFFFLITLIVLITITLYWQTEGKRIIMRL 368
QY      121  HEQIINEGKDKMFLIEKLKIQDMEKANPSSVLERREVEQOGFLHGHDSGLDLRSR 180
Db      369  HEQIINEGKDKMFLIEKLKIQDMEKANPSSVLERREVEQOGFLHGHDSGLDLRSR 428
QY      181  RSVQEGNPRA 190
Db      429  RSVQEGNPRA 438

```

# RESULT 9

AAB83079 standard; Protein; 460 AA.

AAB83079;

29-JUN-2001 (first entry)

Human CASB6411 protein.

Human: CASB6411; vaccine; gene therapy; immunoprophylaxis;  
ovarian cancer; colon cancer; autoimmune disease.

Homo sapiens.

WO200123417-A2.

05-APR-2001.

27-SEP-2000; 2000WO-EP09500.

30-SEP-1999; 99GB-0023154.

07-JUL-2000; 2000GB-0016839.

(SMIK ) SMITHKLINE BEECHAM BIOLOGICALS.

Vinals De Bassols YC;

WPI: 2001-316133/73.

N-PSDB; AAF82460.

Novel CASB6411 polypeptides useful in diagnostics, and as vaccines for  
prophylactic and therapeutic treatment of cancers, particularly ovarian  
and colon cancers, autoimmune diseases and related conditions -  
Claim 1; Page 64; 95pp; English.

The present sequence is human CASB6411 polypeptide. The  
invention relates to CASB6411 polypeptides comprising a sequence  
having at least 70% identity to a sequence of 460 or 154 amino acids  
fully defined in the specification. CASB6411 polypeptides and  
polynucleotides are useful for treating a subject by immunoprophylaxis  
or therapy. The CASB6411 polypeptides are useful in diagnostics, and  
as vaccines for prophylactic and therapeutic treatment of cancers,  
particularly ovarian and colon cancers, autoimmune diseases and related  
conditions. CASB6411 polypeptides are also useful for the  
structure-based design of agonists, antagonists or inhibitors of the  
polypeptide. The full length mRNA encoding the present sequence may  
be alternatively spliced to generate a mRNA encoding a truncated  
CASB6411 protein.

Sequence 460 AA;

Query Match 100.0%; Score 1002; DB 22; Length 460;  
Best Local Similarity 100.0%; Pred. No. 7e-111;  
Matches 190; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1 MNFOPPSKAMRASQMMTEFFILFFPSPFTGVCTLAITITWRLKPSADCGPFGDLPLFIH 60  
|||||



```

|||||
Db 271 MNQPSKARASOMTFEFLFPSTGVLCTLAITWRLKPSADCGPFRGLPIFIH 330
Qy 61 STYSWIDTLSTRPGYLMWVWYIRNLIGSVHFFLTLLVILITLYLWQITGRRIMRL 120
Db 331 STYSWIDTLSTRPGYLMWVWYIRNLIGSVHFFLTLLVILITLYLWQITGRRIMRL 390
Qy 121 HEQIINSGKDKMFLIEKLKIDMEKKANPSSVLERREVEOQGFLLHGEHDSLDLSR 180
Db 391 HEQIINSGKDKMFLIEKLKIDMEKKANPSSVLERREVEOQGFLLHGEHDSLDLSR 450
Qy 181 RSVOEGNPRA 190
Db 451 RSVOEGNPRA 460

RESULT 10
ABP41828
ID ABP41828 standard; Protein: 305 AA.
AC ABP41828;
DX 22-AUG-2002 (first entry)
XX
DE Human ovarian antigen HACMU05, SEQ ID NO:2960.
XX
KW Human; ovarian antigen; ovary; ovarian; breast; cancer; tumour;
KW ovarian cancer; breast cancer; tumour; reproductive system disorder;
KW infertility; pregnancy disorder; anovulation; polycystic ovary syndrome;
KW PCOS; ovarian cyst; dysmenorrhoea; endocrine disorder; infection;
KW inflammatory condition; immune disorder; blood disorder;
KW cardiovascular disorder; respiratory disorder; neurological disorder;
KW gastrointestinal disorder; urinary system disorder; drug screening;
KW gene therapy; chromosome mapping; forensic analysis;
KW antibody preparation; cytostatic; immunomodulatory; neuroprotective;
KW antiinflammatory; gynaecological; reproductive; chromosome 17q25.
XX
OS Homo sapiens.
XX
PN W0200200677-A1.
XX
PD 03-JAN-2002.
XX
PE 07-JUN-2001; 2001MO-US18569.
XX
PR 07-JUN-2000; 2000US-209467P.
XX
PA (HUMA-) HUMAN GENOME SCI INC.
XX
PI Birse CE, Rosen CA;
XX
DR WPI: 2002-147878/19.
DR N-PSDB: ABQ54905.
XX
PT Isolated nucleic acid molecules encoding novel ovarian polypeptides,
PT useful in the prevention, treatment and diagnosis of cancer (e.g.
PT ovarian cancer), immune disorders, cardiovascular disorders and
PT neurological diseases -
XX
PS Claim 11; SEQ ID NO 2960; 2922bp; English.
XX
CC The invention relates to 2175 novel human ovarian antigens (ABP41054-
CC ABP43228) and to cDNAs encoding them (ABQ54131-ABQ56305), and also
CC encompasses polypeptides 90% identical and polynucleotides 95% identical
CC to the sequences of the invention. The invention additionally relates to
CC recombinant vectors and host cells comprising human ovarian antigen
CC polynucleotides, antibodies against human ovarian antigens, and the use
CC of ovarian antigen polynucleotides and polypeptides in diagnosing,
CC treating, prognosing or preventing various ovary and/or breast-related
CC disorders. Such conditions include ovarian cancer and breast cancer, and
CC metastatic tumours of ovarian or breast origin, reproductive system
CC disorders (e.g., infertility, disorders of pregnancy, anovulation,
CC polycystic ovary syndrome, ovarian cysts, and dysmenorrhoea), endocrine

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CC disorders, infections (e.g., chlamydia, HIV, toxoplasmosis, and toxic
CC shock syndrome), inflammatory conditions (e.g., mastitis, oophoritis and
CC vaginitis), immune disorders (e.g., congenital and acquired
CC immunodeficiencies, autoimmune oophoritis, systemic lupus erythematosus),
CC blood-related disorders (e.g., anaemia), cardiovascular disorders,
CC respiratory disorders, neurological disorders, gastrointestinal disorders
CC and urinary system disorders. Ovarian antigen polypeptides and
CC polynucleotides may also be used in screening for compounds which
CC modulate ovarian antigen expression or activity. The polynucleotides may
CC further be used for gene therapy, chromosome mapping, in the
CC identification of individuals and in forensic analysis, and the
CC polypeptides may be used as food additives or to prepare antibodies
CC useful in disease diagnosis, drug targeting and phenotyping. The present
CC sequence represents a human ovarian antigen of the invention.
CC Note: The sequence data for this patent did not form part of the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pcl_sequences.
XX
SQ Sequence 305 AA;
XX
Query Match 29.2%; Score 292.5; DB 23; Length 305;
Best Local Similarity 34.9%; Pred. No. 2.0e-26;
Matches 60; Conservative 36; Mismatches 63; Indels 13; Gaps 2;

Qy 1 MNQPSKARASOMTFEFLFPSTGVLCTLAITWRLKPSADCGPFRGLPIFIH 60
Db 133 LANQARRRPPWPLASHMSTVFLLTLCFPAFLGAAYFLCAVWQVPSSTCGFRLLDIME 192
Qy 61 STYSWIDTL-STRPGYLMWVWYIRNLIGSVHFFLTLLVILITLYLWQITGRRIMRL 119
Db 193 AGRWVWRLLEAGRGVSMLEPWVHRVLYMNTFVFLVSALLAVIYLNQVVRGGRKVICL 252
Qy 120 LHEQIINSGKDKMFLIEKLKIDMEKKANPSSVLERREVEOQGFLLHGEHDSLDLSR 159
Db 253 LKEQISMEGEDIKPLINKLSIVYERKERSRVGTTEAAPALLTDED 304

RESULT 11
ABB39891
ID ABB39891 standard; Peptide: 31 AA.
XX
AC ABB39891;
XX
DX 04-FEB-2002 (first entry)
XX
DE Peptide #7397 encoded by human foetal liver single exon probe.
XX
KW Human; foetal liver; gene expression; single exon nucleic acid probe.
XX
OS Homo sapiens.
XX
PN W0200157277-A2.
XX
PD 09-AUG-2001.
XX
PE 30-JAN-2001; 2001MO-US00669.
XX
PR 04-FEB-2000; 2000US-0180312.
PR 26-MAY-2000; 2000US-0207456.
PR 30-JUN-2000; 2000US-0608408.
PR 03-AUG-2000; 2000US-0632366.
PR 21-SEP-2000; 2000US-0234687.
PR 27-SEP-2000; 2000US-0236359.
PR 04-OCT-2000; 2000GB-0024263.
XX
PA (MOLE-) MOLECULAR DYNAMICS INC.
XX
PI Penn SG, Hanzel DK, Chen W, Rank DR;
XX
DR WPI: 2001-483447/52.
XX
PT Human genome-derived single exon nucleic acid probes useful for
PT analysing gene expression in human fetal liver -

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XX Claim 27; SEQ ID NO 32526; 639pp + sequence listing; English.  
PS  
XX  
CC The invention relates to a single exon nucleic acid probe for  
CC measuring human gene expression in a sample derived from human foetal  
CC liver. The single exon nucleic acid probes may be used for predicting,  
CC measuring and displaying gene expression in samples derived from human  
CC fetal liver. The present sequence is a peptide encoded by a single exon  
CC nucleic acid probe of the invention.  
CC Note: The sequence data for this patent did not form part of the  
CC printed specification, but was obtained in electronic format directly  
CC from WIPO at ftp.wipo.int/pub/published\_pct\_sequences.  
XX  
SQ Sequence 31 AA;  
  
Query Match 14.8%; Score 148; DB 22; Length 31;  
Best Local Similarity 100.0%; Pred. No. 2.6e-10;  
Matches 31; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
  
OY 131 KMFLIEKLIKLDMEKKANPSSIVLERREVE 161  
DB 1 KMFLIEKLIKLDMEKKANPSSIVLERREVE 31  
|||||  
  
RESULT 12  
AAM60631  
ID AAM60631 standard; Protein; 31 AA.  
XX  
AC AAM60631;  
XX  
DT 05-NOV-2001 (first entry)  
XX  
DE Human brain expressed single exon probe encoded protein SEQ ID NO: 32736.  
XX  
DE Human brain expressed single exon probe encoded protein SEQ ID NO: 32736.  
XX  
DE Human; brain expressed exon; gene expression analysis; probe;  
KW microarray; Alzheimer's disease; multiple sclerosis; schizophrenia;  
KW epilepsy; cancer.  
XX  
OS Homo sapiens.  
XX  
PN WO200157275-A2.  
XX  
PD 09-AUG-2001.  
XX  
PF 30-JAN-2001; 2001WO-US00667.  
XX  
PR 04-FEB-2000; 2000US-0180312.  
PR 26-MAY-2000; 2000US-0207456.  
PR 30-JUN-2000; 2000US-0608408.  
PR 03-AUG-2000; 2000US-0632366.  
PR 21-SEP-2000; 2000US-0234687.  
PR 27-SEP-2000; 2000US-0236359.  
PR 04-OCT-2000; 2000GB-0024263.  
XX  
XX (MOLE-) MOLECULAR DYNAMICS INC.  
XX  
PA Penn SG, Hanzel DK, Chen W, Rank DR;  
XX  
PI WPI; 2001-483446/52.  
XX  
DR  
XX  
XX Single exon nucleic acid probes for analyzing gene expression in human  
XX brains -  
XX  
PS Example 4; SEQ ID NO: 32736; 650pp + Sequence Listing; English.  
XX  
CC The present invention provides a number of single exon nucleic acid  
CC probes which are derived from genomic sequences expressed in the human  
CC brain. They can be used to measure gene expression in brain cell samples,  
CC which may enable the diagnosis and improved treatment of nervous system  
CC diseases such as Alzheimer's disease, multiple sclerosis, schizophrenia,  
CC epilepsy and cancers. The present sequence is a protein encoded by one of  
XX the probes of the invention.

SQ Sequence 31 AA;  
  
Query Match 14.8%; Score 148; DB 22; Length 31;  
Best Local Similarity 100.0%; Pred. No. 2.6e-10;  
Matches 31; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
  
OY 131 KMFLIEKLIKLDMEKKANPSSIVLERREVE 161  
DB 1 KMFLIEKLIKLDMEKKANPSSIVLERREVE 31  
|||||  
  
RESULT 13  
AAM73303  
ID AAM73303 standard; Protein; 31 AA.  
XX  
AC AAM73303;  
XX  
DT 06-NOV-2001 (first entry)  
XX  
DE Human bone marrow expressed probe encoded protein SEQ ID NO: 33609.  
XX  
DE Human bone marrow expressed probe encoded protein SEQ ID NO: 33609.  
XX  
DE Human; bone marrow expressed exon; gene expression analysis; probe;  
KW microarray; cancer; leukaemia; lymphoma; myeloma.  
XX  
OS Homo sapiens.  
XX  
PN WO200157276-A2.  
XX  
PD 09-AUG-2001.  
XX  
PF 30-JAN-2001; 2001WO-US00668.  
XX  
PR 04-FEB-2000; 2000US-0180312.  
PR 26-MAY-2000; 2000US-0207456.  
PR 30-JUN-2000; 2000US-0608408.  
PR 03-AUG-2000; 2000US-0632366.  
PR 21-SEP-2000; 2000US-0234687.  
PR 27-SEP-2000; 2000US-0236359.  
PR 04-OCT-2000; 2000GB-0024263.  
XX  
XX (MOLE-) MOLECULAR DYNAMICS INC.  
XX  
PA Penn SG, Hanzel DK, Chen W, Rank DR;  
XX  
PI WPI; 2001-488900/53.  
XX  
DR  
XX  
XX Human genome-derived single exon nucleic acid probes useful for  
XX analyzing gene expression in human bone marrow -  
XX  
PS Example 4; SEQ ID NO: 33609; 658pp + Sequence Listing; English.  
XX  
CC The present invention provides a number of single exon nucleic acid  
CC probes which are derived from genomic sequences expressed in the human  
CC bone marrow. They can be used to measure gene expression in bone marrow  
CC samples, which may enable the improved diagnosis and treatment of cancers  
CC such as lymphoma, leukaemia and myeloma. The present sequence is a  
XX protein encoded by one of the probes of the invention.  
XX  
SQ Sequence 31 AA;  
  
Query Match 14.8%; Score 148; DB 22; Length 31;  
Best Local Similarity 100.0%; Pred. No. 2.6e-10;  
Matches 31; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
  
OY 131 KMFLIEKLIKLDMEKKANPSSIVLERREVE 161  
DB 1 KMFLIEKLIKLDMEKKANPSSIVLERREVE 31  
|||||  
  
RESULT 14  
AAM33503  
ID AAM33503 standard; Protein; 31 AA.  
XX

AC AAM33503;  
 XX  
 XX 17-OCT-2001 (first entry)  
 XX  
 DE Peptide #7540 encoded by probe for measuring placental gene expression.  
 XX  
 XX Probe; microarray; human; placenta; antenatal diagnosis;  
 KW genetic disorder.  
 KW  
 XX Homo sapiens.  
 OS  
 XX WO200157272-A2.  
 PN  
 XX 09-AUG-2001.  
 PD  
 XX 30-JAN-2001; 2001WO-US00663.  
 PE  
 XX 04-FEB-2000; 2000US-0180312.  
 PR 26-MAY-2000; 2000US-0207456.  
 PR 30-JUN-2000; 2000US-0608408.  
 PR 03-AUG-2000; 2000US-0632366.  
 PR 21-SEP-2000; 2000US-0234687.  
 PR 27-SEP-2000; 2000US-0236359.  
 PR 04-OCT-2000; 2000GB-0024263.  
 PR  
 XX (MOLE-) MOLECULAR DYNAMICS INC.  
 PA  
 XX Penn SG, Hanzel DK, Chen W, Rank DR;  
 PI WPI; 2001-488897/53.  
 DR  
 XX Human genome-derived single exon nucleic acid probes useful for  
 PT analyzing gene expression in human placenta -  
 PS Claim 27; SEQ ID No 33772; 654pp; English.  
 PS  
 XX The present invention relates to single exon nucleic acid probes (SENP;  
 CC see A13315-A157546). The present sequence is a peptide encoded by one  
 CC such probe. The probes are useful for producing a microarray for  
 CC predicting, measuring and displaying gene expression in samples derived  
 CC from human placenta. The probes are useful for antenatal diagnosis of  
 CC human genetic disorders.  
 CC  
 XX Sequence 31 AA;  
 SQ  
 Query Match 14.8%; Score 148; DB 22; Length 31;  
 Best Local Similarity 100.0%; Pred. No. 2.6e-10;  
 Matches 31; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 Oy 131 KMFLTEKLTKLQDMERKANPSSLVREKVE 161  
 DB 1 KMFLTEKLTKLQDMERKANPSSLVREKVE 31  
 RESULT 15  
 ABG43154  
 ID ABG43154 standard; Peptide: 31 AA.  
 AC ABG43154;  
 XX  
 XX 19-AUG-2002 (first entry)  
 DT  
 DE Human peptide encoded by genome-derived single exon probe SEQ ID 32819.  
 XX  
 XX Human; single exon probe; asthma; lung cancer; COPD; ILD;  
 KW chronic obstructive pulmonary disease; interstitial lung disease;  
 KW familial idiopathic pulmonary fibrosis; neurofibromatosis;  
 KW tuberous sclerosis; Gaucher's disease; Niemann-Pick disease;  
 KW Hermansky-Pudlak syndrome; sarcoidosis; pulmonary haemosiderosis;  
 KW pulmonary histiocytosis; lymphangioleiomyomatosis; Karagener syndrome;  
 KW pulmonary alveolar proteinosis; fibrocystic pulmonary dysplasia;  
 KW primary ciliary dyskinesia; pulmonary hypertension;  
 KW hyaline membrane disease.

XX  
 OS Homo sapiens.  
 XX  
 XX WO200186003-A2.  
 XX  
 XX 15-NOV-2001.  
 PD  
 XX 30-JAN-2001; 2001WO-US00665.  
 PE  
 XX 04-FEB-2000; 2000US-180312P.  
 PR 26-MAY-2000; 2000US-207456P.  
 PR 30-JUN-2000; 2000US-0608408.  
 PR 03-AUG-2000; 2000US-0632366.  
 PR 21-SEP-2000; 2000US-234687P.  
 PR 27-SEP-2000; 2000US-236359P.  
 PR 04-OCT-2000; 2000GB-0024263.  
 PR  
 XX (MOLE-) MOLECULAR DYNAMICS INC.  
 PA  
 XX Penn SG, Hanzel DK, Chen W, Rank DR;  
 PI WPI; 2002-114183/15.  
 DR  
 XX Spatially-addressable set of single exon nucleic acid probes, used to  
 PT measure gene expression in human lung samples -  
 PS Claim 27; SEQ ID No 32819; 634pp; English.  
 PS  
 XX The invention relates to a spatially-addressable set of single exon  
 CC nucleic acid probes for measuring gene expression in a sample derived  
 CC from human lung comprising single exon nucleic acid probes having one of  
 CC 12614 nucleic acid sequences mentioned in the specification, or their  
 CC complements or the 12387 open reading frames derived from the 12614  
 CC probes. Also included are a microarray comprising the novel set of  
 CC probes; the novel set of probes which hybridize at high stringency to a  
 CC nucleic acid expressed in the human lung; measuring gene expression in a  
 CC sample derived from human lung, comprising (a) contacting the array with  
 CC a collection of detectably labeled nucleic acids derived from human lung  
 CC mRNA, and (b) measuring the label detectably bound to each probe of  
 CC the array; identifying exons in a eukaryotic genome, comprising  
 CC (a) algorithmically predicting at least one exon from genomic sequences  
 CC of the eukaryote; and (b) detecting specific hybridization of detectably  
 CC labeled nucleic acids from eukaryote lung mRNA, to a single exon probe,  
 CC having a fragment identical to the predicted exon, the probe is included  
 CC in the above mentioned microarray; assigning exons to a single gene,  
 CC comprising (a) identifying exons from genomic sequence by the method  
 CC above and (b) measuring the expression of each of the exons in several  
 CC tissues and/or cell types using hybridization to a single exon  
 CC microarrays having a probe with the exon, where a common pattern of  
 CC expression of the exons in the tissues and/or cell types indicates that  
 CC the exons should be assigned to a single gene; a peptide comprising one  
 CC of 12011 sequences, mentioned in the specification, or encoded by the  
 CC probes/open reading frames (ORF). The probes are used for gene  
 CC expression analysis, and for identifying exons in a gene, particularly  
 CC using human lung derived mRNA and for the study of lung diseases  
 CC such as asthma, lung cancer, chronic obstructive pulmonary disease  
 CC (COPD), interstitial lung disease (ILD), familial idiopathic pulmonary  
 CC fibrosis, neurofibromatosis, tuberous sclerosis, Gaucher's disease,  
 CC Niemann-Pick disease, Hermansky-Pudlak syndrome, sarcoidosis, pulmonary  
 CC haemosiderosis, pulmonary histiocytosis, lymphangioleiomyomatosis,  
 CC pulmonary alveolar proteinosis, Karagener syndrome, fibrocystic  
 CC pulmonary dysplasia, primary ciliary dyskinesia, pulmonary hypertension  
 CC and hyaline membrane disease. The present sequence is a peptide/protein  
 CC encoded by a single exon probe of the invention.  
 CC Note: The sequence data for this patent did not form part  
 CC of the printed specification, but was obtained in electronic  
 CC format directly from WIPO at  
 CC ftp.wipo.int/pub/published\_pct\_sequences.  
 XX  
 SQ Sequence 31 AA;  
 Query Match 14.8%; Score 148; DB 23; Length 31;  
 Best Local Similarity 100.0%; Pred. No. 2.6e-10;

Matches 31; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
Qy 131 KMFLIEKLIKIDMEKKANPSSIVLERREYE 161  
|||||  
Db 1 KMFLIEKLIKIDMEKKANPSSIVLERREYE 31

Search completed: November 9, 2002, 04:30:32  
Job time : 81 secs